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6 SEA FILE=HCAPLUS ABB=ON PLU=ON REACTIVE GROUP AND LINKER AND BIOMOL?

=> d ibib abs 126 1-6

L26 ANSWER 1 OF 6 HCAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER:

2003:262085 HCAPLUS

DOCUMENT NUMBER:

138:268074

CODEN: PIXXD2

TITLE:

Method of attachment

INVENTOR(S):

Odedra, Raj; Pickering, Lee

PATENT ASSIGNEE(S):

Amersham Biosciences UK Limited, UK

SOURCE:

PCT Int. Appl., 26 pp.

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT. NO.			, KIND		DATE			APPLICATION NO.				ο.	DATE				
									_		- -						
WO	WO 2003027677			A2 20030403.			WO 2002-GB4369 20020926										
	W:	ΑE,	AG,	AL,	AM,	AT,	AU,	AZ,	BA,	BB,	BG,	BR,	BY,	ΒZ,	CA,	CH,	CN,
		co,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,	GE,	GH,
		GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	ΚE,	KG,	ΚP,	KR,	ΚZ,	LC,	LK,	LR,
		LS,	LT,	LU,	LV,	ΜA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NO,	NZ,	OM,	PH,
		PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK,	SL,	ТJ,	TM,	TN,	TR,	TT,	TZ,
		UΑ,	UG,	US,	UZ,	VC,	VN,	YU,	ZA,	ZM,	ZW,	AM,	ΑZ,	BY,	KG,	ΚZ,	MD,
		RU,	ТJ,	TM							•						
	RW:	GH,	GM,	ΚE,	LS,	ΜW,	ΜZ,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AT,	BE,	BG,
		CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	ΙE,	ΙT,	LU,	MC,	NL,
		PT,	SE,	SK,	TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	GW,	ML,	MR,
		ΝE,	SN,	TD,	TG												

PRIORITY APPLN. INFO.:

GB 2001-23120 A 20010926

OTHER SOURCE(S):

MARPAT 138:268074

AB Disclosed is a method for attaching biomols. to a solid surface and a compn. for prepg. that surface for attachment. The compn. comprises mols. of Formula I and Formula II which are defined as follows: Y - X - Z - R1 Formula I Y'- X' - Z' - R2 Formula II and wherein R1 is a

biomol., a reactive group or a group capable
of forming a reactive group; R2 is different to R1 and
is present in at least a 104 fold molar excess to R1;Y and Y' are groups
which can bind to a solid surface; X and X' are atoms which are, at least,
bivalent; and Z and Z' are linker groups.

L26 ANSWER 2 OF 6 HCAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER:

PATENT ASSIGNEE(S):

2003:42337 HCAPLUS

DOCOME

138:91395

TITLE:

 ${\tt Method} \ \ {\tt for} \ \ {\tt increasing} \ \ {\tt hydrophilicity} \ \ {\tt of} \ \underline{{\tt fluorescent}}$

label compounds, and their use

INVENTOR(S):

Meltola, Niko; Soini, Aleksi Arctic Diagnostics Oy, Finland

SOURCE:

PCT Int. Appl., 46 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE: English FAMILY ACC. NUM. COUNT: PATENT INFORMATION: KIND DATE APPLICATION NO. DATE PATENT NO. _____ _____ WO 2002-FI581 20020701 WO 2003004569 W: AE, AG, AL, AM, AT, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, A1 20030116 CN, CO, CR, CU, CZ, CZ, DE, DE, DK, DK, DM, DZ, EC, EE, EE, ES, FI, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG PRIORITY APPLN. INFO.: FI 2001-1438 A 20010702 US 2001-301831P P 20010702 OTHER SOURCE(S): MARPAT 138:91395 The invention relates to fluorescent label compds. in the form of dipyrrometheneboron difluoride dye derivs. contg. NHCH(CH2CH2Z)CONHY or NHCZCH2CH2CONHY groups, wherein Z is a reactive group and Y is a water-solubilizing moiety or CH2CH2SO3X, with X being a cation. The invention also relates to the use of the compds. in bioanal. assays and cytol. or histol. staining methods. The invention further relates to a method for increasing the hydrophilicity of fluorescent compds. In an example, a glutamic acid-taurine linker, HO2CCH2CH2CH(NH2)CONHCH2CH2SO3H, was prepd. and condensed with 4,4-difluoro-5-(2-thienyl)-1,3-dimethyl-4-bora-3a,4a-diaza-s-indacene-2propionic acid succinimidyl ester and the product was then re-esterified with N-hydroxysuccinimide to give a fluorescent compd. suitable for labeling of mouse IgG anti-AFP. REFERENCE COUNT: THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS g RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L26 ANSWER 3 OF 6 HCAPLUS COPYRIGHT 2003 ACS on STN 1999:126933 HCAPLUS ACCESSION NUMBER: DOCUMENT NUMBER: 130:179636 TITLE: Polysaccharide conjugates of biomolecules and preparation of modified polysaccharides having pendant aldehydes for conjugate preparation Mehta, Harshvardhan; Singh, Rajendra INVENTOR(S): Behringwerke Aktiengesellschaft, Germany PATENT ASSIGNEE(S): SOURCE: PCT Int. Appl., 54 pp. CODEN: PIXXD2 DOCUMENT TYPE: Patent LANGUAGE: English FAMILY ACC. NUM. COUNT: PATENT INFORMATION: PATENT NO. KIND DATE APPLICATION NO. DATE

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PATENT NO. KIND DATE APPLICATION NO. DATE

WO 9907744 Al 19990218 WO 1997-US13803 19970808
W: CA, JP
RW: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE
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19991208 EP 1997-937122 19970808 EP 961783 A1 R: AT, BE, CH, DE, ES, FR, GB, IT, LI, LU, NL, SE JP 1999-512057 JP 2001516394 20010925 19970808 Т2 WO 1997-US13803 W 19970808 PRIORITY APPLN. INFO.: Compds. that are modified polysaccharides having pendant aldehyde functionalities are disclosed. Each of the aldehyde functionalities is attached through a linker to a position corresponding to a hydrogen atom of a different hydroxyl group of unmodified polysaccharide. Also disclosed is a method for introducing an amine-reactive functionality into a dextran. The method comprises (a) reacting the dextran with an alkylating agent having a functionality that reacts with an hydroxyl group of the dextran thereby forming an alkylated dextran wherein the alkylating agent has an olefin group and (b) treating the alkylated dextran to convert the olefin group to an amine-reactive functionality. A polysaccharide can be conjugated to a biomol. by carrying out the above method and reacting the amine-reactive functionality with an amine functionality on the biomol. to produce polysaccharide conjugated to the biomol. Protein A-dextran conjugate was prepd. by reacting dextran with allyl glycidyl ether, subjecting the product to ozonolysis, and reacting with protein A. The conjugate was used in an assay to detect autoantibodies to the 65 kD isoform of glutamic acid decarboxylase (GAD65) in human serum samples, esp. from patients with insulin dependent diabetes mellitus. REFERENCE COUNT: 3. THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L26 ANSWER 4 OF 6 HCAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER:

1997:542488 HCAPLUS

DOCUMENT NUMBER:

127:206935

TITLE:

Benzophenoxazine dyes for labeling of

biomolecules

INVENTOR(S):

Simmonds, Adrian; Miller, James N.; Moody, Christopher John; Swann, Elizabeth; Briggs, Mark Samuel Jonathan;

Bruce, Ian Edward

PATENT ASSIGNEE(S):

Amersham International PLC, UK; Simmonds, Adrian; Miller, James N.; Moody, Christopher John; Swann, Elizabeth; Briggs, Mark Samuel Jonathan; Bruce, Ian

Edward

SOURCE:

PCT Int. Appl., 39 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 3

PATENT INFORMATION:

PATENT NO.				KIND DATE				APPLICATION NO.					ο.	DATE				
₩ 0 9729154			A1 19970814				WO 1997-GB324					19970205						
	W:	AL,	AM,	AT,	AU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	CA,	CH,	CN,	CU,	CZ,	DE,	
		DK,	EE,	ES,	FI,	GB,	GE,	HU,	IL,	IS,	JP,	ΚE,	KG,	ΚP,	KR,	ΚZ,	LC,	
		LK,	LR,	LS,	LT,	LU,	LV,	MD,	MG,	MK,	MN,	MW,	MX,	NO,	NZ,	PL,	PT,	
		RO,	RU,	SD,	SE,	SG,	SI,	SK,	ТJ,	TM,	TR,	TT,	UA,	ŬĠ,	US,	UZ,	VN,	
		ΥU,	AM,	ΑZ,	BY,	KG,	ΚZ,	MD,	RU,	ТJ,	TM							
	RW:	ΚE,	LS,	MW,	SD,	SZ,	UG,	ΑT,	BE,	CH,	DE,	DK,	ES,	FI,	FR,	GB,	GR,	
		ΙE,	IT,	LU,	MC,	NL,	PT,	SE,	BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	ML,	
		MR,	ΝE,	SN,	TD,	TG												
EP	8852	63		A	1 :	1998:	1223		E	P 199	97-9	0247	7	1997	0205			

EP:885263 В1 20010425 R: CH, DE, DK, FR, GB, IT, LI, NL, SE JP 2000504755 Т2 20000418 JP 1997-528276 19970205 19970828 AU 1997-16115 AU 9716115 **A1** 19970206 US 6166202 Α 20001226 US 1999-117608 19990115

PRIORITY APPLN. INFO.: GB 1996-2265 19960205 Α WO 1997-GB324 19970205 W

OTHER SOURCE(S): MARPAT 127:206935

GΙ

AB Benzophenoxazine compds. having formula I (X = O, NH, N-alkyl, N-aryl, N-alkenyl; Y = H, NR1R2; R1, R2 = C1-12 alkyl, aryl, alkenyl, LA; L =C0-20 linker which may contain O, N, S; A = amino, amide, CN, hydroxy, thiol, carboxy, sulfonate, phosphate, a reactive group by means of which the compd. can be linked to a biomol.) are synthesized. The compds. are used as fluorescent dyes for labeling biomols.

L26 ANSWER 5 OF 6 HCAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 1997:34059 HCAPLUS

DOCUMENT NUMBER: 126:57117

TITLE: Methods for the production of platinum-based

linkers between labels and bio-organic

molecules, for labeling bio-organic molecules, for detecting biological substances of interest and

diagnostic test kits

Houthoff, Hendrik Jan; Reedijk, Jan; Jelsma, Tinka; INVENTOR(S):

Van Es, Remco Maria; Van Den Berg, Franciscus Michiel; Lempers, Edwin Leo Mario; Bloemink, Marieke Johanna

PATENT ASSIGNEE(S):

Kreatech Biotechnology B.V., Neth.; Houthoff, Hendrik Jan; Reedijk, Jan; Jelsma, Tinka; Van Es, Remco Maria;

Van Den Berg, Franciscus Michiel; Lempers, Edwin Leo

Mario; Bloemink, Marieke Johanna

PCT Int. Appl., 36 pp. SOURCE:

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE ----WO 9635696 **A**1 19961114 WO 1996-NL198 19960508 AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT,

LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN CA 2218815 19961114 CA 1996-2218815 19960508 AA AU 9657040 19961129 AU 1996-57040 A1

AU 724320 B2 20000914 JP 11505533 Т2 19990521 JP 1996-533965 19960508 NZ 307633 20000128 NZ 1996-307633 19960508 Α EP 1019420 A1 20000719 EP 1996-915218 19960508

EP 1019420 20030806 В1

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI

PRIORITY APPLN. INFO.:

EP 1995-201197 A 19950509 WO 1996-NL198 W 19960508

OTHER SOURCE(S): CASREACT 126:57117; MARPAT 126:57117

The present invention provides improved methods of producing platinum compds., which are very suitable for producing labeled substances, which can be used to detect specific mols. of interest. The platinum coordination compds. have two reactive groups of which one is replaced by a label and the other one can be replaced by a substance to be labeled. Prodn. of labeled substances is very much improved by selection of the right starting materials and producing the right intermediates. The efficiency of labeling is very much improved, thereby enabling the prodn. of labeling kits which are also a part of the present invention. The methods can be used for the detection of, e.g., various microorganisms and gene translocations/abnormalities.

L26 ANSWER 6 OF 6 HCAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NÜMBER:

1987:583601 HCAPLUS

DOCUMENT NUMBER:

107:183601

TITLE:

Macrocyclic bifunctional chelating agents

INVENTOR(S):

Meares, Claude F.; DeNardo, Sally J.; Cole, William

C.; Mol, Min K.

PATENT ASSIGNEE(S):

University of California, Berkeley, USA

SOURCE:

U.S., 11 pp.

CODEN: USXXAM

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 4678667	Α	19870707	US 1985-751769	19850702
PRIORITY APPLN. INFO.	:		US 1985-751769	19850702
~~				

$$R_1$$
 $N-(CH_2)_1-N$ R_2 $L-(CH_2)_k$ $(CH_2)_m$ $N-(CH_2)_n-N$

AB A Cu chelate conjugate for diagnostic or therapeutic applications which involve Cu2+ localization via the systemic route comprises a Cu2+ chelate of a bifunctional macrocyclic polyamide I (k, l, m, n = 1-4; .gtoreq.2 ofR1-R4 = -CH2CO2-, the rest H; L = linker with chem. reactive group capable of reacting with a biomol ., and CH2(1) indicates that the ring C to which L is attached has only 1 H) and, chem. linked to the linker, a biomol. selected from among antibodies, antibody fragments, serum proteins, and bleomycin. The Cu2+ chelate of 13-(p-nitrobenzyl)-1,4,8,11-tetraazacyclotetradecane-N,N',N'',N'''-tetraacetic acid (prepn. given) was activated by conversion of the arom. NO2 group to a bromoacetamide by the method of Meares, et al. (1984), then mixed (1.5 mM) with Lym-1 antibody (IgG2 mouse monoclonal antibody against B-cell lymphoma) (0.5 mM) in 0.15 M Na3PO4 and reacted for 2 h at 37.degree. and pH 9.0-9.5 (adjusted with satd. Na3PO4). The resulting antibody-chelate product was purified on a Sephadex G-50-80 column.